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Tuofu Zhu, David Muthui, Sarah Holte, David Nickle, Feng Feng, Scott Brodie, Yon Hwangbo, James I. Mullins, and Lawrence Corey

Evidence for Human Immunodeficiency Virus Type 1 Replication In Vivo in CD14+ Monocytes and Its Potential Role as a Source of Virus in Patients on Highly Active **Antiretroviral Therapy** 

J. Virol. 2002 76: 707-716. [Abstract] [Full Text] [PDF]

🗂 Guislaine Carcelain, Roland Tubiana, Assia Samri, Vincent Calvez, Constance Delaugerre, Henri Agut, Christine Katlama, and Brigitte Autran

Transient Mobilization of Human Immunodeficiency Virus (HIV)-Specific CD4 T-Helper Cells Fails To Control Virus Rebounds during Intermittent Antiretroviral Therapy in Chronic HIV Type 1 Infection

J. Virol. 2001 75: 234-241. [Abstract] [Full Text] [PDF]

Monika Hermankova, Janet D. Siliciano, Yan Zhou, Daphne Monie, Karen Chadwick, Joseph B. Margolick, Thomas C. Quinn, and Robert F. Siliciano

Analysis of Human Immunodeficiency Virus Type 1 Gene Expression in Latently Infected Resting CD4<sup>+</sup> T Lymph cytes In Vivo

J. Virol. 2003 77: 7383-7392. [Abstract] [Full Text] [PDF]

Wei Lu and Jean-Marie Andrieu

In Vitro Human Immunodeficiency Virus Eradicati n by Autol g us CD8+ T Cells Expanded with Inactivated-Virus-Pulsed Dendritic Cells

J. Virol. 2001 75: 8949-8956. [Abstract] [Full Text] [PDF]
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Mark Dybul, Marybeth Daucher, Mark A. Jensen, Claire W. Hallahan, Tae-Wook Chun, Michael Belson, Bertha Hidalgo, David C. Nickle, Christian Yoder, Julia A. Metcalf, Richard T. Davey, Linda Ehler, Diane Kress-Rock, Elizabeth Nies-Kraske, Shuying Liu, James I. Mullins, and Anthony S.
Genetic Characterization of Rebounding Human Immunodeficiency Virus Type 1 in Plasma during Multiple Interruptions of Highly Active Antiretroviral Therapy J. Virol. 2003 77: 3229-3237. [Abstract] [Full Text] [PDF]
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L5	L3 and interferon\$ with Hiv\$2	204	L5
L4	(interferon-alpha or interferon adj alpha) with (2a or 2b) same HIV-1	6	L4
L3	(interferon adj alpha\$3 or interferon-alpha\$3 or interferon adj alpha adj (2a or 2b)) same HIV\$2	293	L3
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L2 L3 L4 L5 L6 L7 L8 L9 L10 L11 L12 L13 L14 L15 L16 L17 L18 L19 L20	01 JUL 2003  4062 S (INTERFERON ADJ ALPHA OR INF-ALPHA OR INTERFERON-ALPHA OR INF 2223 S (INTERFERON ADJ ALPHA OR INF-ALPHA OR INTERFERON-ALPHA OR INF 2909 S (INTERFERON (A) ALPHA OR INF-ALPHA OR INTERFERON-ALPHA OR INF 990 S L3 AND (CTL OR NK OR LYMPHOCYTE OR T (A) CELL OR T-CELL) 50 S L3 AND (PEG OR PEGYLAT####) 15 S L5 AND L4 7 DUP REM L6 (8 DUPLICATES REMOVED) 77 S L3 AND (HAART OR HIGHLY (A) ACTIVE (A) (ANTIRETROVIRAL OR AN 43 DUP REM L8 (34 DUPLICATES REMOVED) 12 S L9 AND L5 10 S L10 NOT L7 2563 S L3 NOT PY>2000 856 S L12 AND L4 8 S L12 AND L5 7 DUP REM L14 (1 DUPLICATE REMOVED) 4 S L15 NOT (L11 OR L7) 3458 S (INTERFERON (3N) ALPHA) (S) (HIV OR AIDS OR IMMUNODEFICIENCY 54 S L17 AND (PEG OR PEGYLAT####) 37 DUP REM L18 (17 DUPLICATES REMOVED) 17 S L19 NOT (L7 OR L11 OR L16)
L2 L3 L4 L5 L6 L7 L8 L9 L10 L11 L12 L13 L14 L15 L16 L17 L18 L19 L20 L21	01 JUL 2003  4062 S (INTERFERON ADJ ALPHA OR INF-ALPHA OR INTERFERON-ALPHA OR INF 2223 S (INTERFERON ADJ ALPHA OR INF-ALPHA OR INTERFERON-ALPHA OR INF 2909 S (INTERFERON (A) ALPHA OR INF-ALPHA OR INTERFERON-ALPHA OR INF 990 S L3 AND (CTL OR NK OR LYMPHOCYTE OR T (A) CELL OR T-CELL) 50 S L3 AND (PEG OR PEGYLAT####) 15 S L5 AND L4 7 DUP REM L6 (8 DUPLICATES REMOVED) 77 S L3 AND (HAART OR HIGHLY (A) ACTIVE (A) (ANTIRETROVIRAL OR AN 43 DUP REM L8 (34 DUPLICATES REMOVED) 12 S L9 AND L5 10 S L10 NOT L7 2563 S L3 NOT PY>2000 856 S L12 AND L4 8 S L12,AND L5 7 DUP REM L14 (1 DUPLICATE REMOVED) 4 S L15 NOT (L11 OR L7) 3458 S (INTERFERON (3N) ALPHA) (S) (HIV OR AIDS OR IMMUNODEFICIENCY 54 S L17 AND (PEG OR PEGYLAT####) 37 DUP REM L18 (17 DUPLICATES REMOVED) 17 S L19 NOT (L7 OR L11 OR L16) 10773 S (INTERFERON) (S) (HIV OR AIDS OR IMMUNODEFICIENCY OR HIV-1)
L2 L3 L4 L5 L6 L7 L8 L9 L10 L11 L12 L13 L14 L15 L16 L17 L18 L19 L20 L21 L22	01 JUL 2003  4062 S (INTERFERON ADJ ALPHA OR INF-ALPHA OR INTERFERON-ALPHA OR INF 2223 S (INTERFERON ADJ ALPHA OR INF-ALPHA OR INTERFERON-ALPHA OR INF 2909 S (INTERFERON (A) ALPHA OR INF-ALPHA OR INTERFERON-ALPHA OR INF 990 S L3 AND (CTL OR NK OR LYMPHOCYTE OR T (A) CELL OR T-CELL) 50 S L3 AND (PEG OR PEGYLAT###) 15 S L5 AND L4 7 DUP REM L6 (8 DUPLICATES REMOVED) 77 S L3 AND (HAART OR HIGHLY (A) ACTIVE (A) (ANTIRETROVIRAL OR AN 43 DUP REM L8 (34 DUPLICATES REMOVED) 12 S L9 AND L5 10 S L10 NOT L7 2563 S L3 NOT PY>2000 856 S L12 AND L4 8 S L12,AND L5 7 DUP REM L14 (1 DUPLICATE REMOVED) 4 S L15 NOT (L11 OR L7) 3458 S (INTERFERON (3N) ALPHA) (S) (HIV OR AIDS OR IMMUNODEFICIENCY 54 S L17 AND (PEG OR PEGYLAT####) 37 DUP REM L18 (17 DUPLICATES REMOVED) 17 S L19 NOT (L7 OR L11 OR L16) 10773 S (INTERFERON) (S) (HIV OR AIDS OR IMMUNODEFICIENCY OR HIV-1) 5279 S L21 AND (PEG OR PEGYLAT#### OR CTL OR NK OR LYMPHOCYTE OR T
L2 L3 L4 L5 L6 L7 L8 L9 L10 L11 L12 L13 L14 L15 L16 L17 L18 L19 L20 L21	01 JUL 2003  4062 S (INTERFERON ADJ ALPHA OR INF-ALPHA OR INTERFERON-ALPHA OR INF 2223 S (INTERFERON ADJ ALPHA OR INF-ALPHA OR INTERFERON-ALPHA OR INF 2909 S (INTERFERON (A) ALPHA OR INF-ALPHA OR INTERFERON-ALPHA OR INF 990 S L3 AND (CTL OR NK OR LYMPHOCYTE OR T (A) CELL OR T-CELL) 50 S L3 AND (PEG OR PEGYLAT####) 15 S L5 AND L4 7 DUP REM L6 (8 DUPLICATES REMOVED) 77 S L3 AND (HAART OR HIGHLY (A) ACTIVE (A) (ANTIRETROVIRAL OR AN 43 DUP REM L8 (34 DUPLICATES REMOVED) 12 S L9 AND L5 10 S L10 NOT L7 2563 S L3 NOT PY>2000 856 S L12 AND L5 7 DUP REM L14 (1 DUPLICATE REMOVED) 4 S L15 NOT (L11 OR L7) 3458 S (INTERFERON (3N) ALPHA) (S) (HIV OR AIDS OR IMMUNODEFICIENCY 54 S L17 AND (PEG OR PEGYLAT####) 37 DUP REM L18 (17 DUPLICATES REMOVED) 17 S L19 NOT (L7 OR L11 OR L16) 10773 S (INTERFERON) (S) (HIV OR AIDS OR IMMUNODEFICIENCY OR HIV-1) 5279 S L21 AND (PEG OR PEGYLAT#### OR CTL OR NK OR LYMPHOCYTE OR T 3991 S L22 NOT PY>2000 701 S L23 AND INTERFERON (S) (COMBINATION OR HAART OR ANTIVIRAL OR
L2 L3 L4 L5 L6 L7 L8 L9 L10 L11 L12 L13 L14 L15 L16 L17 L18 L19 L20 L21 L22 L23	4062 S (INTERFERON ADJ ALPHA OR INF-ALPHA OR INTERFERON-ALPHA OR INTER 2223 S (INTERFERON ADJ ALPHA OR INF-ALPHA OR INTERFERON-ALPHA OR INF 2909 S (INTERFERON (A) ALPHA OR INF-ALPHA OR INTERFERON-ALPHA OR INF 990 S L3 AND (CTL OR NK OR LYMPHOCYTE OR T (A) CELL OR T-CELL) 50 S L3 AND (PEG OR PEGYLAT####) 15 S L5 AND L4 7 DUP REM L6 (8 DUPLICATES REMOVED) 77 S L3 AND (HAART OR HIGHLY (A) ACTIVE (A) (ANTIRETROVIRAL OR AN 43 DUP REM L8 (34 DUPLICATES REMOVED) 12 S L9 AND L5 10 S L10 NOT L7 2563 S L3 NOT PY>2000 856 S L12 AND L5 7 DUP REM L14 (1 DUPLICATE REMOVED) 4 S L5 NOT (L11 OR L7) 3458 S (INTERFERON (3N) ALPHA) (S) (HIV OR AIDS OR IMMUNODEFICIENCY 54 S L17 AND (PEG OR PEGYLAT####) 37 DUP REM L18 (17 DUPLICATES REMOVED) 17 S L19 NOT (L7 OR L11 OR L16) 10773 S (INTERFERON) (S) (HIV OR AIDS OR IMMUNODEFICIENCY OR HIV-1) 5279 S L21 AND (PEG OR PEGYLAT#### OR CTL OR NK OR LYMPHOCYTE OR T 3991 S L22 NOT PY>2000 701 S L23 AND INTERFERON (S) (COMBINATION OR HAART OR ANTIVIRAL OR 340 DUP REM L24 (361 DUPLICATES REMOVED)

L7 ANSWER 1 OF 7 MEDLINE DUPLICATE 1

AN 2003086386 MEDLINE

DN 22485890 PubMed ID: 12598770

TI Perforin expression in **T cells** and virological response to **PEG**-interferon alpha2b in HIV-1 infection.

- AU Portales Pierre; Reynes Jacques; Rouzier-Panis Regine; Baillat Vincent; Clot Jacques; Corbeau Pierre
- CS Laboratoire d'Immunologie, Hopital Saint Eloi, Service des Maladies Infectieuses et Tropicales, Hopital Gui de Chauliac, Montpellier, France.
- SO AIDS, (2003 Mar 7) 17 (4) 505-11. Journal code: 8710219. ISSN: 0269-9370.
- CY England: United Kingdom
- DT Journal; Article; (JOURNAL ARTICLE)
- LA English
- FS Priority Journals; AIDS
- EM 200304
- ED Entered STN: 20030225

  Last Updated on STN: 20030430

  Entered Medline: 20030439
- Entered Medline: 20030429 AB OBJECTIVE AND DESIGN: Interferon alpha (IFNalpha), which is known to directly inhibit the HIV-1 replicative cycle and to increase the activity of cytotoxic T lymphocytes (CTL), is being tested as an anti-HIV agent. As CTL play a major role in immune defence against HIV, we wanted to further characterize CTL activity and the effect of IFNalpha on it. METHODS: We followed by flow cytometry the intracellular expression of the key mediator of cytotoxicity, perforin, in peripheral blood T cells of patients treated with IFNalpha. RESULTS: We observed that the percentage of T cells harbouring perforin was higher in infected subjects than in non-infected controls. Administration of IFNalpha2b attached to polyethylene glycol increased this perforin expression further and reduced viral load (P = 0.010). The increase in the percentage of **T** cells expressing perforin correlated with IFNalpha-induced decrease in viral load (r, 0.753; P = 0.003). In addition, the level of perforin expression before IFNalpha administration was inversely correlated with viral load remaining after IFNalpha administration (r, -0.647; P= 0.017). CONCLUSION: The pre-therapeutic percentage of perforin-positive T cells might be a predictive marker of the virological response to IFNalpha in HIV-1-infected patients.
- L7 ANSWER 2 OF 7 MEDLINE

DUPLICATE 2

- AN 2003086385 MEDLINE
- DN 22485889 PubMed ID: 12598769
- TI Interferon-alpha restores HIV-induced alteration of natural killer cell perforin expression in vivo.
- AU Portales Pierre; Reynes Jacques; Pinet Valerie; Rouzier-Panis Regine; Baillat Vincent; Clot Jacques; Corbeau Pierre
- CS Laboratoire d'Immunologie, Hopital Saint Eloi, the Service des Maladies Infectieuses et Tropicales, Hopital Gui de Chauliac, Montpellier, France.
- SO AIDS, (2003 Mar 7) 17 (4) 495-504. Journal code: 8710219. ISSN: 0269-9370.
- CY England: United Kingdom
- DT Journal; Article; (JOURNAL ARTICLE)
- LA English
- FS Priority Journals; AIDS
- EM 200304
- ED Entered STN: 20030225

Last Updated on STN: 20030430 Entered Medline: 20030429

) cells are known to be decreased in HIV-infected patients. However, the mechanisms responsible for this NK deficiency are poorly understood. Because of the role of NK cells in the host defence against microbial infections, this defect contributes to the virus-induced immune deficiency. The aim of the present study was to better understand this defect in order to be able to restore NK function in HIV infection. DESIGN AND METHODS: The expression of the cytolytic mediators perforin and granzyme A was analysed by flow cytometry, the lytic activity of peripheral blood NK cells of HIV-infected patients was analysed by cytotoxic assay, and the expression of perforin was followed during administration of interferon (IFN)alpha attached to polyethylene glycol (PEG)-IFNalpha. RESULTS: The lytic activity and the expression of perforin and granzyme A was low in NK cells of infected individuals in comparison with normal control volunteers. both groups NK cytotoxic capacity was linked to perforin expression. The low perforin expression in HIV-infected subjects negatively correlated with HIV RNA plasma level. administration of PEG-IFNalpha restored perforin expression even in patients whose viral load was not reduced by this treatment. CONCLUSIONS: These results suggest that  ${\tt HIV}{ ext{-}}{\tt induced}$   ${\tt NK}$  deficiency could be partly mediated by a defect in perforin and granzyme A expression, and that PEG -IFNalpha could be used in infected subjects to directly improve their natural immunity in addition to eventually reducing their viraemia. ANSWER 3 OF 7 CAPLUS COPYRIGHT 2003 ACS L7 2001:676618 CAPLUS AN DN135:225873 HIV-specific immune response promoted by interferon-. ΤI alpha. IN Laughlin, Mark A. PA Schering Corporation, USA SO PCT Int. Appl., 34 pp. CODEN: PIXXD2 DT Patent LΑ English FAN.CNT 1 APPLICATION NO. DATE PATENT NO. KIND DATE \_\_\_\_\_\_ \_ \_ \_ \_ \_\_\_\_\_ WO 2001-US7453 20010308 PΙ WO 2001066132 A2 20010913 WO 2001066132 Α3 20020124 W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, HR, HU, ID, IL, IN, IS, JP, KG, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD,  $\mathtt{MG}$ ,  $\mathtt{MK}$ ,  $\mathtt{MN}$ ,  $\mathtt{MZ}$ ,  $\mathtt{NO}$ ,  $\mathtt{NZ}$ ,  $\mathtt{PL}$ ,  $\mathtt{PT}$ ,  $\mathtt{RO}$ ,  $\mathtt{RU}$ ,  $\mathtt{SE}$ ,  $\mathtt{SG}$ ,  $\mathtt{SI}$ ,  $\mathtt{SK}$ ,  $\mathtt{SL}$ , TJ, TM, TR, TT, TZ, UA, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG US 2002004584 A1 20020110 US 2001-801980 20010308 EP 1263457 A2 20021211 EP 2001-922303 20010308 AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR BR 2001008997 20030603 BR 2001-8997 20010308 Α NO 2002004255 NO 2002-4255 20020906 20020906 Α PRAI US 2000-188338P 20000309 Ρ WO 2001-US7453 W 20010308 AB Use of interferon-.alpha., e.g., pegylated

interferon .alpha.-2a or 2b for prepn. of a medicament for promotion of an HIV-1 specific immune response,

OBJECTIVE: The percentage and the activity of natural killer (NK

AΒ

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e.g., promotion of HIV-1 specific T-
     cells, in adult and pediatric patients having HIV-
     1 infections as well as patients co-infected with HIV-
     1 and HCV comprising a therapeutically effective amt. of
     pegylated interferon-.alpha., e.g.,
     pegylated interferon .alpha.-2b is disclosed.
     ANSWER 4 OF 7 CAPLUS COPYRIGHT 2003 ACS
L7
ΑN
     2001:265599 CAPLUS
DN
     134:294522
TI
     Interferon .alpha. homologues
     Heinrichs, Volker; Chen, Teddy; Patten, Phillip A.
IN
PA
     Maxygen, Inc., USA
SO
     PCT Int. Appl., 209 pp.
     CODEN: PIXXD2
DT
     Patent
T.A
     English
FAN.CNT 1
                                            APPLICATION NO.
                      KIND DATE
     PATENT NO.
                       A2
                            20010412
                                            WO 2000-US27781 20001006
PΙ
     WO 2001025438
                      A3
                            20020711
     WO 2001025438
             AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
             CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR,
             HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT,
             LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU,
             SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
         RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY,
             DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ,
             CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
                                                             20001006
                                            EP 2000-970665
                           20020911
     EP 1238082
                       A2
             AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
             IE, SI, LT, LV, FI, RO, MK, CY, AL
                                            JP 2001-528590
                                                             20001006
                       T2
                            20030325
     JP 2003511031
                             19991007
PRAI US 1999-415183
                       Α
                             20001006
     WO 2000-US27781
                       W
     .alpha. Interferon homologues (both nucleic acids and polypeptides) are
AB
     provided. Compns. including these interferon homolog polypeptides and
     nucleic acids, recombinant cells comprising said homolog polypeptides and
     nucleic acids, methods of making the new homologues, antibodies to the new
     homologues, and methods of using the homologues are provided. Integrated
     systems comprising the sequences of the nucleic acids or polypeptides are
     also provided. These interferon .alpha. homologues are useful for
     inhibiting tumor growth and viral replication, and are also useful for
     treating autoimmune diseases.
     ANSWER 5 OF 7 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC. DUPLICATE 3
L7
AN
     2001:530219 BIOSIS
DN
     PREV200100530219
     Effect of HCV viral dynamics on treatment design: Lessons learned from
ΤI
     HIV.
     Bain, Vincent G. (1)
AU
     (1) Department of Medicine, Division of Gastroenterology, University of
CS
     Alberta, 8440 112 Street, 2E1.14 WMC, Edmonton, AB, T6G 2R7 Canada
     American Journal of Gastroenterology, (October, 2001) Vol. 96, No. 10, pp.
SO
     2818-2828. print.
     ISSN: 0002-9270.
     Article; General Review
DT
     English
LΑ
SL
     English
```

Viral load measurements provide an indication of viral replication, and AB thereby serve as a valuable tool to guide the initiation of therapy and subsequent changes. Plasma human immunodeficiency viral load strongly predicts the rate of decrease in CD4+ lymphocyte count, and progression to AIDS and death. Furthermore, the efficacy of antiretroviral therapy can be assessed by monitoring changes in plasma human immunodeficiency viral load. Similarly, viral load provides valuable information about the natural history of the hepatitis C virus infection. Hepatitis C viral load can be used to predict the likelihood of response to standard interferon-alpha treatment and other interferon-alpha regimens and to monitor treatment efficacy. Increased understanding of the natural history of the hepatitis C virus infection and the nature of resistance to interferon-alpha therapy suggests that effective treatment regimens must maintain serum levels of interferonalpha. Ideally, interferon-alpha serum levels should provide constant pressure on the virus and should prevent viral rebound, thereby avoiding continued viral replication and minimizing the potential for emergence of resistant quasi-species. Current regimens designed to address these points include early aggressive intervention, combination drug regimens, prolonged maintenance, and novel interferons. By enabling the design and rapid assessment of new treatment regimens, viral load measurement will revolutionize the clinical management of the hepatitis C virus infection, as it has the HIV.

- L7 ANSWER 6 OF 7 MEDLINE
- AN 2001457575 MEDLINE
- DN 21395681 PubMed ID: 11504966
- TI Early control of HIV replication in primary HIV-1 infection treated with antiretroviral drugs and **pegylated** IFN alpha: results from the Primoferon A (ANRS 086) Study.
- AU Emilie D; Burgard M; Lascoux-Combe C; Laughlin M; Krzysiek R; Pignon C; Rudent A; Molina J M; Livrozet J M; Souala F; Chene G; Grangeot-Keros L; Galanaud P; Sereni D; Rouzioux C
- CS Service de Medecine Interne et d'Immunologie Clinique, Hopital Antoine Beclere, Institut Paris-Sud sur les Cytokines, Clamart, France. (Primoferon A Study Group).
- SO AIDS, (2001 Jul 27) 15 (11) 1435-7. Journal code: 8710219. ISSN: 0269-9370.
- CY England: United Kingdom
- DT (CLINICAL TRIAL)

  Journal; Article; (JOURNAL ARTICLE)

  (MULTICENTER STUDY)
- LA English
- FS Priority Journals; AIDS
- EM 200109
- ED Entered STN: 20010816 Last Updated on STN: 20010924 Entered Medline: 20010920
- AB IFN alpha has both antiviral and immunostimulating properties. The ANRS086 Primoferon A Study evaluated in 12 patients with primary HIV infection the tolerance and efficacy of an early and transient administration of pegylated IFN alpha, in addition to highly active antiretroviral therapy. Tolerance was good, and this regimen allowed the early control of HIV replication and rapid decay of the viral reservoir. These results support the initiation of comparative studies with pegylated INF alpha in primary HIV infection.
- L7 ANSWER 7 OF 7 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.
- AN 2002:153875 BIOSIS

- DN PREV200200153875
- TI Immunologic profile of HIV-1 infected patients after administration of pegylated interferon-alpha -2b (PEG-intron.
- AU Vigklis, V. (1); Laughlin, M.; Gargalianos, P.; Lazanas, M.; Botsi, C.; Saroglou, G.; Sabatakou, H.; Paraskeva, D.; Stavrianeas, N.; Mavroidi, N.; Glue, P.; Hatzakis, A. (1)
- CS (1) Dept. of Hygiene and Epidemiology, Athens Univ. Medical School, Athens Greece
- SO AIDS (Hagerstown), (October, 2000) Vol. 14, No. Supplement 4, pp. S19. http://www.aidsonline.com/. print.
  Meeting Info.: Fifth International Congress on Drug Therapy in HIV Infection Glasgow, UK October 22-26, 2000 ISSN: 0269-9370.
- DT Conference
- LA English

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L11 ANSWER 1 OF 10 MEDLINE
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- AN 2002098072 MEDLINE
- DN 21686113 PubMed ID: 11827840
- TI Interferon-alpha in treatment of chronic hepatitis C in co-infected HIV-patients in combination with ribavirin and as a pre-load therapy in treatment-naive HIV-positive patients. 8th European Conference on Clinical Aspects and Treatment of HIV Infection (8th ECCATH), 29-31 October, Athens Greece.
- AU Tossing Gudrun
- CS ESSEX Pharma GmbH Muenchen, Tannenstr. 15a, D-42653 Solingen, Germany.. qudrun.tossing@essex.de
- SO EUROPEAN JOURNAL OF MEDICAL RESEARCH, (2002 Jan 29) 7 (1) 44-6. Journal code: 9517857. ISSN: 0949-2321.
- CY Germany: Germany, Federal Republic of
- DT Conference; Conference Article; (CONGRESSES)
- LA English
- FS Priority Journals
- EM 200204
- ED Entered STN: 20020206

Last Updated on STN: 20020423 Entered Medline: 20020422

AB Though AIDS-related morbidity and mortality are generally decreasing as a result of highly active antiretroviral

therapy (HAART) and prevention of opportunistic

infections, dual infection with HCV and HIV leads to an acceleration in the natural course of chronic hepatitis C (cHC) and worsening of associated liver disease and complications. Mortality from co-morbid HCV infection within this population is increasing and has become a major challenge in the management of HIV-related complications. As treatment strategies to fight cHC have been essentially ameliorated within the recent two years in using pegylated interferon-alfa2b ( Peg-IFN-alfa2b) combined with ribavirin, t here is hope that the successful therapeutic outcomes in HCV-mono-infected individuals may be partly translated into benefits for the difficult-to-treat patients with HCV-HIV co-infection. A number of issues arise when beginning HCV treatment during HAART, as for instance possible interactions with antiretroviral therapies, increased risk of special side effects, and a compromise in adherence due to the addition of new medication in patients already taking several drugs. On the other hand there is also the chance that Peg-IFN-alfa2b fights HIV as well as HCV. First data of pre-load therapy with Peg-IFN-alfa2b in treatment-na ve HIV-positive individuals before the initiation of HAART have also been presented during the 8th European Conference on Clinical Aspects and Treatment of HIV Infection (8th ECCATH), October 2001 in Athens.

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L11 ANSWER 2 OF 10 CAPLUS COPYRIGHT 2003 ACS
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AN 2002:928015 CAPLUS

DN 137:379981

TI HIV therapy

IN Laughlin, Mark A.; Glue, Paul W.; Stalgis, Carlos O.

PA USA

SO U.S. Pat. Appl. Publ., 14 pp.

CODEN: USXXCO

DT Patent

LA English

FAN.CNT 1

PATENT NO. KIND DATE APPLICATION NO. DATE

PI US 2002182179 A1 20021205 US 2000-516673 20000301

PRAI US 1999-122370P P 19990302

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US 1999-128296P
                      P
                            19990408
AΒ
     Methods for the treatment of treatment-naive as well as
     treatment-experienced adult and pediatric patients with HIV-1 infections
     as well as patients co-infected with HIV-1 and HCV involving
     administration of a therapeutically effective amt. of pegylated
     interferon-alfa, e.g., pegylated interferon alfa-2b as
     monotherapy or preferably in assocn. with a therapeutically effective amt.
     of at least one of ribavirin, IL-2, IL-12, pentafuside alone or in
     combination with a therapeutically effective amt. of an anti-HIV-1 drug
     therapy, e.g., HAART are disclosed.
L11 ANSWER 3 OF 10 CAPLUS COPYRIGHT 2003 ACS
AN
     2000:790325 CAPLUS
DN
     133:329566
     PEGylated interferon- alpha - CCR5 antagonist
ΤI
     combination HIV therapy
IN
     Laughlin, Mark A.
     Schering Corporation, USA
PA
     PCT Int. Appl., 80 pp.
SO
     CODEN: PIXXD2
DT
     Patent
LΑ
     English
FAN.CNT 1
                      KIND DATE
                                           APPLICATION NO.
     PATENT NO.
                                                            DATE
                                           _____
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     WO 2000066141
                                           WO 2000-US11634 20000501
                      A2
                            20001109
PΤ
                      A3
                            20010208
     WO 2000066141
         W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR,
             CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, HR, HU, ID, IL, IN,
             IS, JP, KG, KR, KZ, LC, LK, LR, LT, LU, LV, MA, MD, MG, MK, MN,
             MX, NO, NZ, PL, PT, RO, RU, SE, SG, SI, SK, SL, TJ, TM, TR, TT,
             TZ, UA, US, UZ, VN, YU, ZA, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
         RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE,
             DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF,
             CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
     EP 1175224
                      A2
                                          EP 2000-928604
                          20020130
                                                            20000501
             AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
             IE, SI, LT, LV, FI, RO
     BR 2000010593
                                           BR 2000-10593
                                                            20000501
                       Α
                            20020213
     JP 2002543144
                       T2
                            20021217
                                           JP 2000-615025
                                                            20000501
                                           NO 2001-5367
     NO 2001005367
                       Α
                            20020103
                                                            20011102
PRAI US 1999-304897
                            19990504
                       Α2
                            20000501
     WO 2000-US11634
                       W
OS
     MARPAT 133:329566
AB
     The invention discloses the use of a PEGylated
     interferon-.alpha. and a CCR5 antagonist, further in
     assocn. with at least one of ribavirin, IL-2, IL-12, pentafuside alone or
     in combination with an anti-HIV-1 drug therapy, e.g.,
     HAART (highly active antiretroviral
     therapy), for prepn. of a medicament for the treatment of
     HIV-1 infections as well as HIV-1
     infections and HCV co-infections in treatment-naive as well as
     treatment-experienced adult and pediatric patients.
L11 ANSWER 4 OF 10 CAPLUS COPYRIGHT 2003 ACS
     2000:628016 CAPLUS
AN
DN
     133:206775
     HIV therapy using pegylated interferon-alfa alone and in assocn.
TI
     with anti-HIV-1 drug therapy
     Laughlin, Mark A.; Glue, Paul W.; Stalgis, Carlos O.
IN
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US 1999-124304P

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19990312

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PCT Int. Appl., 45 pp.
SO
     CODEN: PIXXD2
DT
     Patent
     English
LΑ
FAN.CNT 1
                      KIND DATE
                                            APPLICATION NO.
                                                             DATE
     PATENT NO.
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                      A2
ΡI
                            20000908
                                            WO 2000-US5361
                                                             20000301
     WO 2000051631
                      A3
                            20010118
     WO 2000051631
            AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CZ,
             DE, DK, DM, EE, ES, FI, GB, GD, GE, HR, HU, ID, IL, IN, IS, JP, KG, KR, KZ, LC, LK, LR, LT, LU, LV, MA, MD, MG, MK, MN, MX, NO,
             NZ, PL, PT, RO, RU, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA,
             US, UZ, VN, YU, ZA, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
         RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE,
             DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF,
             CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
                            20000919
                                           JP 2000-55695
                                                              20000301
     JP 2000256211
                       A2
                                            EP 2000-301695
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                            20000913
     EP 1034790
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     EP 1034790
                            20001213
                       Α3
         R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
             IE, SI, LT, LV, FI, RO
                       A2
PRAI US 1999-260388
                            19990302
                       Α2
                            19990312
     US 1999-268521
                       A2
     US 1999-288358
                            19990408
     US 1999-454004
                       A2
                            19991203
     The uses of pegylated interferon-alfa, alone, and in assocn.
AB
     with an anti-HIV-1 drug therapy, and ribavirin for the prepn. of a
     medicament for treating treatment-naive as well as treatment-experienced
     adult and pediatric patients having HIV-1 infections as well as patients
     co-infected with HIV-1 and hepatitis C virus (HCV) involving comprising a
     therapeutically effective amt. of pegylated interferon-alfa,
     e.g., pegylated interferon alfa-2b as monotherapy or preferably
     in assocn. with a therapeutically effective amt. of at least one of
     ribavirin, IL-2, IL-12, pentafuside alone or in combination with a
     therapeutically effective amt. of an anti-HIV-1 drug therapy, e.g.,
     HAART are disclosed.
     ANSWER 5 OF 10 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.
L11
AN
     2003:236115 BIOSIS
DN
     PREV200300236115
     Treatment of chronic hepatitis C in HIV infected patients (PTS)
ΤI
     with pegylated interferon alpha 2B (
     PEG-IFN) plus ribavirin (RBV) versus pegylated
     interferon alpha 2B.
ΑU
     Cargnel, A. (1); Angeli, E. (1); Mainini, A. (1); Casella, A. (1);
     Gubertini, G. (1); Giorgi, R. (1); Orlando, G. (1); Duca, P. G.
     (1) II Dept Infectious Diseases, L. Sacco Hospital, Milan, Italy Italy
CS
     Journal of Hepatology, (April 2003, 2003) Vol. 38, No. Supplement 2, pp.
SO
     132. print.
     Meeting Info.: 38th Annual Meeting of the European Association for the
     Study of the Liver Istanbul, Turkey March 29-April 01, 2003 European
     Association for the Study of the Liver
     . ISSN: 0168-8278.
DT
     Conference
     English
LА
     ANSWER 6 OF 10 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.
L11
     2002:618390 BIOSIS
AN
     PREV200200618390
DN
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PA

Schering Corporation, USA

- ΤI An open, multicenter, randomized trial comparing pegylated interferon alpha-2B (PEG-IFN) plus ribavirin (RBV) versus PEG-IFN for treatment of HIV/HCV co-infected patients.
- Carquel, Antonietta (1); Angeli, Elena (1); Casella, Alberto (1); ΑU Gubertini, Guido (1); Mainini, Annalisa (1); Orlando, Giovanna (1); Duca, Piergiorgio
- (1) II Dept Infect Diseases, Sacco Hospital, Milan Italy CS
- Hepatology, (October, 2002) Vol. 36, No. 4 Part 2, pp. 363A. SO http://hepatology.aasldjournals.org/scripts/om.dll/serve?action=searchDB&s earchDBfor=home&id=jhep. print. Meeting Info.: 53rd Annual Meeting on the Liver BOSTON, MA, USA November 01-05, 2002 ISSN: 0270-9139.
- DTConference
- LΑ English
- ANSWER 7 OF 10 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC. L11
- AN 2002:566005 BIOSIS
- PREV200200566005 DN
- ΤI Early response to combination therapy with low-dose pegylated interferon-alpha (Peg-IFN) 2b and ribavirin for chronic hepatitis C in HIV-infected patients.
- Moreno, L. (1); Quereda, C. (1); Moreno, A. (1); Casado, J. L. (1); AU Dronda, F. (1); Perez-Elias, M. J. (1); Mateos, M. L. (1); Moreno, S. (1)
- (1) Ramon y Cajal Hospital, Madrid Spain
- Abstracts of the Interscience Conference on Antimicrobial Agents and Chemotherapy, (2001) Vol. 41, pp. 285. print. Meeting Info.: 41st Annual Meeting of the Interscience Conference on Antimicrobial Agents and Chemotherapy Chicago, Illinois, USA September 22-25, 2001
- DTConference
- LΑ
- English AΒ Background: Response to conventional interferon-alpha has been poor in HIV patients (pts) with chronic hepatitis C (hepC). We evaluate the safety and efficacy of Peg-IFN and ribavirin (RBV) combined therapy in the treatment of chronic hepC in pts coinfected with HIV. Methods: Open, prospective, ongoing study. Pts included had coinfection with HCV and HIV, persistently elevated ALT levels, and a liver biopsy showing either portal or bridging fibrosis. Therapy include Peg-IFN-alpha 2b (50 mcg weekly) in combination with RBV (800 mg b.i.d) during 12 months. Primary endpoint is sustained virological response (no detectable HCV RNA by PCR) 6 months after end of therapy. Results: 35 pts were included. Baseline characteristics: median HIV RNA levels and CD4 count were 1.7 log and 544 cells/mm3. 32 pts were on antiretroviral (ARV) therapy (AZT:11; d4T:21 pts). Median ALT and AST values were 83 and 61, respectively, and median HCV RNA was 9X105 IU/mL (7X104-8X106). Most frequent genotypes include 1a (8), 1b (10), 2a (2), and 3a (11). Fibrosis score was 1,8 (1-3). Evaluation at 24 weeks (n=18 pts). 6 (33.3%) patients are HCV RNA negative, and 10 (55%) patients have normalized transaminases levels. Response at 3rd month predicted response at 6th month (66% vs 8%, p=0,02). Toxic side effects were: influenza-like-illness (88%), tiredness (82%), local inflammatory reaction (28%), headache (20%), citopenies (17%), depression (14%). Treatment had to be discontinued in 5 pts (14%). No apparent interaction in toxicity or efficacy has been observed in pts who are on HAART. Median CD4 count has declined to 347 cells/mm3 (264-595). HIV RNA has been maintained or decreased in all the pts. Conclusions: Low dose Peg-IFN-alpha in combination with RBV is safe in HIV pts with chronic hepC, including those who receive ARV drugs. A poor response (33%) has been observed at week 24.

This result suggests the need of use of higher doses in these pts.

ANSWER 8 OF 10 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC. L112002:520492 BIOSIS ANPREV200200520492 DNAntiretroviral efficacy of pegylated-alpha interferon. TIMoreno, L. (1); Quereda, C. (1); Moreno, M. E. (1); Moreno, A. (1); ΑU Gutierrez, C. (1); Marti-Belda, P. (1); Diz, S. (1); Moreno, S. (1) (1) Ramon y Cajal Hospital, Madrid Spain CS SO Abstracts of the Interscience Conference on Antimicrobial Agents and Chemotherapy, (2001) Vol. 41, pp. 347. print. Meeting Info.: 41st Annual Meeting of the Interscience Conference on Antimicrobial Agents and Chemotherapy Chicago, Illinois, USA September 22-25, 2001 Conference DT LΑ English Background: It has been suggested that the new formulation of AΒ alpha-interferon, the pegylated alpha -interferon, recently introduced for the treatment of chronic hepatitis C, may have a clinically significant antiretroviral activity. Methods: Prospective, open clinical trial of therapy of chronic hepatitis C in **HIV**-infected subjects with a combination of low-dose pegylated interferon (50 micrograms once weekly) and ribavirin (800 mg/d). HIV RNA was measured at months 1, 3 and 6. Results: Among  $\overline{3}5$  patients included in the study, 23 had  ${\tt HIV}$  RNA <50 copies/mL and 12 had detectable  ${\tt HIV}$  RNA. All the patients with undetectable viral load were on highly active antiretroviral therapy (HAART) and none developed virological rebound during the 6 months of therapy with pegylated interferon. Among the patients with detectable viral load, 3 were naive for antiretroviral therapy and 9, who had developed virological failure, were on stable therapy. The median CD4 in the 12 patients was 551 cells/mm3 (350-729) and the median HIV RNA was 3 log copies/mL (1.8-3.6). 75% of patients, after one month of treatment with pegylated interferon and ribavirin, had a median viral load decreased of 0.6 log (0.1 to 1.2). In the third month of treatment, the decrease of median viral load had remained of 0.5 log. Two-thirds of the patients had a sustained response through the period of treatment, without modifications of their antiretroviral therapy. Conclusions: Therapy for chronic hepatitis C with a combination of pegylated interferon and ribavirin may help control HIV viremia both in naive and pretreated subject. Future clinical trials should define the role of pegylated interferon in the treatment of HIV infection. ANSWER 9 OF 10 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC. L11 AN 2002:153874 BIOSIS DN PREV200200153874 HIV-1 dynamics in vivo after administration of ΤI pegylated interferon-alpha-2b (PEG -intron. Sypsa, V. (1); Laughlin, M.; Gargalianos, P.; Lazanas, M.; Botsi, C.; ΑU Saroglou, G.; Anastassopoulou, C. (1); Sabatakou, H.; Magafas, N.; Stavrianeas, N.; Giannakopoulou, P.; Glue, P.; Hatzakis, A. (1) (1) Dept of Hygiene and Epidemiology, Athens Univ. Medical School, Athens CS AIDS (Hagerstown), (October, 2000) Vol. 14, No. Supplement 4, pp. S18. SO http://www.aidsonline.com/. print. Meeting Info.: Fifth International Congress on Drug Therapy in HIV Infection Glasgow, UK October 22-26, 2000

DT Conference

ISSN: 0269-9370.

#### LA English

- L11 ANSWER 10 OF 10 EMBASE COPYRIGHT 2003 ELSEVIER SCI. B.V.
- AN 2000129506 EMBASE
- TI [Coinfection with the hepatitis C virus and HIV: Current aspects]. CO-INFECTION PAR LE VIRUS DE L'HEPATITE C ET LE VIRUS DE L'IMMUNODEFICIENCE HUMAINE: ASPECTS ACTUELS.
- AU Bani-Sadr F.; Perronne C.
- CS F. Bani-Sadr, Svc. Maladies Infectieuses/Tropic., Hop. Universitaire Raymond-Poincare, Faculte de Medecine Paris-Ouest, 104, boulevard Raymond-Poincare, 92380 Garches, France
- SO Medecine et Maladies Infectieuses, (2000) 30/SUPPL. 1 (43-48). Refs: 39

ISSN: 0399-077X CODEN: MMAIB5

- CY France
- DT Journal; Conference Article
- FS 004 Microbiology 037 Drug Literature Index 048 Gastroenterology
- LA French
- SL English; French
- AB The treatment of coinfection with the hepatitis C virus (HCV) in HIV- infected patients was rarely discussed before the era of the HIV protease inhibitors, since the response to monotherapy with interferon alpha (INF.alpha.) was

poor, with a mean prognosis of the **HIV** disease estimated at around ten years. In the present context, monitoring is reconsidered. The **HIV**-associated immunosuppression may be responsible for a false negativity of some serologic tests for HCV. The **HIV**-HCV coinfection increases the risk of maternofoetal transmission of HCV. Studies evaluating the influence of the **HIV** coinfection on the natural history of the HCV infection show its deleterious role. The immune restoration obtained with the **highly active** 

antiretroviral therapies is not linked with a decrease of the HCV vital load. The HIV-HCV coinfection is responsible for a threefold increase of the risk of elevation of seric transaminases when an antiretroviral treatment is given. The immune restoration obtained with an antiretroviral treatment may reveal the HCV infection and favor a rapid aggravation of hepatic histology and evolution toward cirrhosis. HCV-associated complications may become a major factor of morbidity and mortality, leading to the need for an anti-hepatitis C treatment in HIV-infected patients. The combination of INF.

alpha. and ribavirin seems to be the best treatment. Its efficacy and tolerability must be evaluated in HIV-infected patients. Drug interactions are likely to occur, and INF.alpha.,

like ribavirin, may favor CD4 lymphopenia. A new form of INF.

alpha. with a prolonged half-life (PEG-INF.

alpha.) seems to be promising. (C) 2000 Editions scientifiques et medicales Elsevier SAS.

- L16 ANSWER 1 OF 4 SCISEARCH COPYRIGHT 2003 THOMSON ISI
- AN 2001:496512 SCISEARCH
- GA The Genuine Article (R) Number: 442BH
- TI New therapies for the treatment of AIDS-related Kaposi sarcoma
- AU Dezube B J (Reprint)
- CS Beth Israel Deaconess Med Ctr, Div Hematol Oncol, 330 Brookline Ave, CC-913, Boston, MA 02215 USA (Reprint); Beth Israel Deaconess Med Ctr, Div Hematol Oncol, Boston, MA 02215 USA
- CYA USA
- SO CURRENT OPINION IN ONCOLOGY, (SEP 2000) Vol. 12, No. 5, pp. 445-449. Publisher: LIPPINCOTT WILLIAMS & WILKINS, 530 WALNUT ST, PHILADELPHIA, PA 19106-3621 USA. ISSN: 1040-8746.
- DT General Review; Journal
- LA English
- REC Reference Count: 34
  - \*ABSTRACT IS AVAILABLE IN THE ALL AND IALL FORMATS\*
- AB Kaposi sarcoma (KS) is the most common tumor arising in HIV -infected patients and is an AIDS-defining illness by the Centers for Disease Control guidelines. The clinical course of AIDS-related KS is highly variable, ranging from minimal stable disease to explosive growth. Recent advances in the elucidation of the pathogenesis of KS are uncovering many potential targets for KS therapies. Such targets include the processes of angiogenesis and cellular differentiation, sex hormones, and the KS herpesvirus/human herpesvirus-8. With the increasing recognition that effective antiretroviral regimens are associated with both a decreased proportion of new AIDS-defining KS cases and a regression in the size of existing KS lesions, most, if not all, KS patients should be advised to take antiretroviral drugs that will maximally decrease HIV-1 viral load. Five agents are currently approved by the Food and Drug Administration for the treatment of KS: alitretinoin gel for topical administration; and liposomal daunorubicin, liposomal doxorubicin, paclitaxel, and interferonalpha for systemic administration. Many more agents, particularly angiogenesis inhibitors, are in early clinical development. The potential interaction between anti-KS agents and antiretroviral agents needs to be kept in mind. Virtually all patients with KS can derive benefit from the many approved and investigational agents developed through years of collaborative translational and clinical research. Curr Opin Oncol 2000. 12:445-449 (C) 2000 Lippincott Williams & Wilkins, Inc.
- L16 ANSWER 2 OF 4 SCISEARCH COPYRIGHT 2003 THOMSON ISI
- AN 2001:30994 SCISEARCH
- GA The Genuine Article (R) Number: 371RQ
- TI Immunologic profile of HIV-1 infected patients after administration of pegylated interferon-alpha -2b (PEG-intron)
- AU Vigklis V (Reprint); Laughlin M; Gargalianos P; Lazanas M; Botsi C; Saroglou G; Sabatakou H; Paraskeva D; Stavrianeas N; Mavroidi N; Glue P; Hatzakis A
- CS Univ Athens, Sch Med, Dept Hyg & Epidemiol, Athens, Greece; Schering Plough Res Inst, Kenilworth, NJ USA; G Gennimates Hosp, Athens, Greece; Red Cross Hosp, Athens, Greece; Syggrou Hosp, Athens, Greece; Evangelismos Hosp, Athens, Greece
- CYA Greece; USA
- SO AIDS, (OCT 2000) Vol. 14, Supp. [4], pp. S19-S19. MA P8.
  Publisher: LIPPINCOTT WILLIAMS & WILKINS, 530 WALNUT ST, PHILADELPHIA, PA
  19106-3621 USA.
  ISSN: 0269-9370.
- DT Conference; Journal

LA English

REC Reference Count: 0

L16 ANSWER 3 OF 4 SCISEARCH COPYRIGHT 2003 THOMSON ISI

AN 2001:30993 SCISEARCH

- GA The Genuine Article (R) Number: 371RQ
- TI HIV-1 dynamics in vivo after administration of pegylated interferon-alpha-2b(PEG -intron)
- AU Sypsa V (Reprint); Laughlin M; Gargalianos P; Lazanas M; Botsi C; Saroglou G; Anastassopoulou C; Sabatakou H; Magafas N; Stavrianeas N; Giannakopoulou P; Glue P; Hatzakis A
- CS Univ Athens, Sch Med, Dept Hyg & Epidemiol, Athens, Greece; Schering Plough Res Inst, Kenilworth, NJ USA; G Gennimatas Hosp, Pireas, Greece; Red Cross Hosp, Pireas, Greece; Syggrou Hosp, Athens, Greece; Evangelismos Hosp, Athens, Greece

CYA Greece; USA

- SO AIDS, (OCT 2000) Vol. 14, Supp. [4], pp. S18-S18. MA P7. Publisher: LIPPINCOTT WILLIAMS & WILKINS, 530 WALNUT ST, PHILADELPHIA, PA 19106-3621 USA. ISSN: 0269-9370.
- DT Conference; Journal

LA English

- REC Reference Count: 1
- L16 ANSWER 4 OF 4 SCISEARCH COPYRIGHT 2003 THOMSON ISI

AN 2001:30992 SCISEARCH

- GA The Genuine Article (R) Number: 371RQ
- TI Safety, tolerability and antiviral pharmacodynamics of pegylated interferon-alpha 2b in HIV-1 infection: a phase I study
- AU Hatzakis A (Reprint); Laughlin M; Gargalianos P; Lazanas M; Botsi C; Saroglou G; Glue P
- CS Univ Athens, Sch Med, Dept Hyg & Epidemiol, Athens, Greece; Schering Plough Res Inst, Kenilworth, NJ USA; G Gennimates Hosp, Pireas, Greece; Tzanion Hosp, Pireas, Greece; Syggrou Hosp, Athens, Greece; Evangelismos Hosp, Athens, Greece

CYA Greece; USA

- SO AIDS, (OCT 2000) Vol. 14, Supp. [4], pp. S18-S18. MA P6.
  Publisher: LIPPINCOTT WILLIAMS & WILKINS, 530 WALNUT ST, PHILADELPHIA, PA
  19106-3621 USA.
  ISSN: 0269-9370.
- DT Conference; Journal

LA English

REC Reference Count: 0

- L20 ANSWER 1 OF 17 MEDLINE
- Development of pegylated interferons for the treatment of ΤI chronic hepatitis C.
- Kozlowski A; Charles S A; Harris J M ΑU
- BioDrugs, (2001) 15 (7) 419-29. Ref: 46 SO Journal code: 9705305. ISSN: 1173-8804.
- L20 ANSWER 2 OF 17 MEDLINE
- TI [Chronic hepatitis C and HIV. Current therapeutic options]. Chronische Hepatitis C und HIV. Aktuelle therapeutische Optionen.
- AII Mauss S
- MMW FORTSCHRITTE DER MEDIZIN, (2001 Apr 2) 143 Suppl 1 46-9. SO Journal code: 100893959. ISSN: 1438-3276.
- ANSWER 3 OF 17 CAPLUS COPYRIGHT 2003 ACS
- Treatment of hepatitis C and anemia in human immunodeficiency ΤI virus-infected patients
- Dieterich, Douglas T. ΑU
- Journal of Infectious Diseases (2002), 185(Suppl. 2), S128-S137 SO CODEN: JIDIAQ; ISSN: 0022-1899
- ANSWER 4 OF 17 CAPLUS COPYRIGHT 2003 ACS L20
- Early control of HIV replication in primary HIV-1 infection treated with TIantiretroviral drugs and pegylated IFN.alpha.: Results from the primoferon A (ANRS 086) study
- Fenton, Kevin A.; Chinouya, Martha; Davidson, Oliver; Copas, Andrew ΑU
- AIDS (London, United Kingdom) (2001), 15(11), 1435-1437 SO CODEN: AIDSET; ISSN: 0269-9370
- L20 ANSWER 5 OF 17 CAPLUS COPYRIGHT 2003 ACS
- PEG-modified lysozyme C and medicinal compositions for the TItreatment of some serious diseases
- Veronesi, Paolo Alberto; Rodriguez, Pablo E. A. IN
- PCT Int. Appl., 33 pp. SO CODEN: PIXXD2
- L20 ANSWER 6 OF 17 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.
- ΤI Hepatitis C and liver fibrosis.
- Schuppan, D. (1); Krebs, A.; Bauer, M.; Hahn, E. G. ΑU
- Cell Death and Differentiation, (January 2003, 2003) Vol. 10, No. SO Supplement 1, pp. S59-S67. print. ISSN: 1350-9047.
- ANSWER 7 OF 17 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC. L20
- ΤI A randomized controlled trial of pegylated-interferon alpha-2b+ribavirin vs interferon alpha -2b+ribavirin in **HIV** co-infected patients.
- Pol, S. (1); Perronne, C. (1); Carrat, F. (1); Banisadr, F. (1); Morand, ΑU P. (1); Lunel, F. (1); Rosenthal, E. (1)
- Journal of Hepatology, (April 2003, 2003) Vol. 38, No. Supplement 2, pp. SO 32. print. Meeting Info.: 38th Annual Meeting of the European Association for the
  - Study of the Liver Istanbul, Turkey March 29-April 01, 2003 European Association for the Study of the Liver . ISSN: 0168-8278.
- L20 ANSWER 8 OF 17 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.
- Pegylated interferon alpha in patients TТ treated from primary HIV infection prevents viral rebound during structured treatment interruptions of antiretroviral drugs.

- AU Emilie, D. (1); Burgard, M.; Krzysiek, R. (1); Rouzious, C.; Galanaud, P. (1)
- SO Journal of Interferon and Cytokine Research, (2002) Vol. 22, No. Supplement 1, pp. S-80. print.

  Meeting Info.: Joint Meeting of the International Society for Interferon and Cytokine Research, the International Cytokine Society, the Society for Leukocyte Biology, and the European Cytokine Society on Cytokines and Interferons Turin, Italy October 06-10, 2002 International Society for Interferon and Cytokine Research

  . ISSN: 1079-9907.
- L20 ANSWER 9 OF 17 EMBASE COPYRIGHT 2003 ELSEVIER SCI. B.V.
- TI Hepatitis C virus in human immunodeficiency virus-infected individuals: An emerging comorbidity with significant implications.
- AU Gonzalez S.A.; Talal A.H.
- SO Seminars in Liver Disease, (2003) 23/2 (149-166).

Refs: 149

ISSN: 0272-8087 CODEN: SLDIEE

- L20 ANSWER 10 OF 17 EMBASE COPYRIGHT 2003 ELSEVIER SCI. B.V.
- TI Peginterferon-.alpha.-2a (40kD) plus ribavirin: A review of its use in the management of chronic hepatitis C.
- AU Keating G.M.; Curran M.P.
- SO Drugs, (2003) 63/7 (701-730).

Refs: 89

ISSN: 0012-6667 CODEN: DRUGAY

- L20 ANSWER 11 OF 17 EMBASE COPYRIGHT 2003 ELSEVIER SCI. B.V.
- TI [Treatment of hepatitis C and B virus infection with interferon].
  PLACE DES INTERFERONS DANS LE TRAITEMENT DES INFECTIONS PAR LES VIRUS DE
  L'HEPATITE B ET DE L'HEPATITE C.
- AU Cacoub P.; Benhamou Y.
- SO Revue de Medecine Interne, (1 Nov 2002) 23/SUPPL. 4 (459s-474s). Refs: 163

ISSN: 0248-8663 CODEN: RMEIDE

- L20 ANSWER 12 OF 17 EMBASE COPYRIGHT 2003 ELSEVIER SCI. B.V.
- TI Potential of interferon-.alpha. in solid tumours: Part 2.
- AU Santhanam S.; Decatris M.; O'Byrne K.
- SO BioDrugs, (2002) 16/5 (349-372).

Refs: 391

ISSN: 1173-8804 CODEN: BIDRF4

- L20 ANSWER 13 OF 17 EMBASE COPYRIGHT 2003 ELSEVIER SCI. B.V.
- TI The natural history and antiviral treatment of hepatitis C in haemophilia.
- AU Lee C.; Dusheiko G.
- SO Haemophilia, (2002) 8/3 (322-329).

Refs: 33

ISSN: 1351-8216 CODEN: HAEMF4

- L20 ANSWER 14 OF 17 EMBASE COPYRIGHT 2003 ELSEVIER SCI. B.V.
- TI Clinical experiences with interferon as monotherapy or in combination with ribavirin in patients co-infected with HIV and HCV.
- AU Puoti M.; Zanni B.; Bruno R.; Airoldi M.; Rossi S.; Roldan E.Q.; El Hamad I.; Moretti F.; Castelli F.; Sacchi P.; Filice G.; Carosi G.
- SO HIV Clinical Trials, (2002) 3/4 (324-332).

Refs: 76

ISSN: 1528-4336 CODEN: HCTIA8

- L20 ANSWER 15 OF 17 SCISEARCH COPYRIGHT 2003 THOMSON ISI
- TI Effects of HCV treatment on HIV course. Preliminary results of

the, anrs hc 02-ribavic study: A randomized, controlled trial of pegylated interferon-alpha 2b with ribavirin vs interferon-alpha 2b with ribavirin for the treatment of chronic HCV in HIV co-infection.

- AU Banisadr F (Reprint); Carrat F; Pol S; Rosenthal E; Morand P; Lunel F; Perronne C
- SO HEPATOLOGY, (OCT 2002) Vol. 36, No. 4, Part 2, Supp. [S], pp. 585A-585A.
  MA 1688.
  Publisher: W B SAUNDERS CO, INDEPENDENCE SQUARE WEST CURTIS CENTER, STE 300, PHILADELPHIA, PA 19106-3399 USA.

ISSN: 0270-9139.

- L20 ANSWER 16 OF 17 SCISEARCH COPYRIGHT 2003 THOMSON ISI
- TI An open, multicenter, randomized trial comparing pegylated interferon alpha-2b (PEG-IFN) plus ribavirin (RBV) versus PEG-IFN for treatment of HIV/HCV co-infected patients.
- AU Cargnel A (Reprint); Angeli E; Casella A; Gubertini G; Mainini A; Orlando G; Duca P
- SO HEPATOLOGY, (OCT 2002) Vol. 36, No. 4, Part 2, Supp. [S], pp. 363A-363A. MA 802.
  Publisher: W B SAUNDERS CO, INDEPENDENCE SQUARE WEST CURTIS CENTER, STE 300, PHILADELPHIA, PA 19106-3399 USA.
  ISSN: 0270-9139.
- L20 ANSWER 17 OF 17 SCISEARCH COPYRIGHT 2003 THOMSON ISI
- TI Hepatitis C and human immunodeficiency virus coinfections
- AU Dodig M; Tavill A S (Reprint)
- SO JOURNAL OF CLINICAL GASTROENTEROLOGY, (NOV-DEC 2001) Vol. 33, No. 5, pp. 367-374.

Publisher: LIPPINCOTT WILLIAMS & WILKINS, 530 WALNUT ST, PHILADELPHIA, PA 19106-3621 USA.

ISSN: 0192-0790.

- L27 ANSWER 1 OF 73 MEDLINE
- TI Antiretrovirals plus immunomodulators: didanosine/interferon-alpha combination shows promise.
- AU Folkers G
- SO NIAID AIDS AGENDA, (1996 Jun) 8-9. Journal code: 9432911.
- L27 ANSWER 2 OF 73 MEDLINE
- TI Immune-based therapies.
- AU Lein B
- SO PI PERSPECTIVE, (1995 Dec) (no 17) 16-8. Journal code: 9102818. ISSN: 1058-7454.
- L27 ANSWER 3 OF 73 MEDLINE
- TI Safety and efficacy of interferon-ribavirin combination therapy in HCV-HIV coinfected subjects: an early report.
- AU Zylberberg H; Benhamou Y; Lagneaux J L; Landau A; Chaix M L; Fontaine H; Bochet M; Poynard T; Katlama C; Pialoux G; Brechot C; Pol S
- SO GUT, (2000 Nov) 47 (5) 694-7. Journal code: 2985108R. ISSN: 0017-5749.
- L27 ANSWER 4 OF 73 MEDLINE
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